

FDAMA STAKEHOLDER MEETING APRIL 28, 1999

Talking with Stakeholders About FDA Modernization

**Your question/comments will become part of Docket Number: 99N-0386 Fax to: 1-888-361-4011 (on April 28 only)

Dr. Mr. Last Name (required)	
Mrs. Ms. John DSTrander	
Organization Warlowide Clinical Trials	
<u>Stakeholder Group</u> ✓ stakeholder group you represent	
☐ Consumer Group ☐ Health Professional ☐ Industry ☐ A	ssociation • Other
<u>Center</u> ✓ the center/product area your comments address	
☐ Center for Biologics ☐ Center for Devices and Radiological Health ☐ Center for Veterinary Medicine ☐ FDA General ☐ Center for Drug Evaluatio ☐ Center for Food Safety and ☐ Office of Regulatory Affair	d Applied Nutrition
Questions to Stakeholders	
Please check the box next to the stakeholder question/s from the March 22, 1999, F question/comment addresses.	ederal Register notice which your
☐ 1. What actions do you propose the Agency take to expand FDA's capability science into its risk-based decision-making?	to incorporate state-of-the-art
□ 2. What actions do you propose to facilitate the exchange and integration of scientific information to better enable FDA to meet its public health responsibilities throughout a product's life cycle?	
3. What actions do you propose for educating the public about the concept of balancing risks against benefits in public health decision-making?	
What actions do you propose to enable FDA and its product centers to for risk to the public health?	ous resources on areas of greatest
What additional actions do you propose for enhancing communication profeedback and/or evaluation of our modernization efforts?	ocesses that allow for ongoing
□ 6. Additional Comments on FDA Modernization Efforts.	
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